Final programme

Thursday, 23 May

13:00-16:00 **Registration**

17:00-17:30 **Welcome coffee**

17:30-20:00 **Opening ceremony**

Greeting of the Guests

'*For HAE Patients*' Award Prize-giving ceremony

Opening lectures I-II *Chairs: Anete Grumach, Marco Cicardi*

I-1 Bradykinin receptors: agonists, antagonists, expression, signaling and adaptation to sustained stimulation
Francois Marceau

I-2 The protease storm of angioedema
Coen Maas

20:00 **Welcome Dinner**

Friday, 24 May

08:30-09:10 **Opening lectures III** *Chairs: Christian Drouet, Péter Gál*

I-3 Activation and regulation of the lectin pathway of the complement system
Péter Gál

09:10-10:25 Scientific lectures
NON-CONVENTIONAL DIAGNOSTICS

O-01 Development of a functional assay for diagnosis of types I and II HAE based on inhibition of factor XIIa or kallikrein
Joseph Kusumam, Baby G. Tholanikunnel, Sonia Bains, Konrad Bork, Anette Bygum, Anne Aabom, Claus Koch, Berhane Ghebrehiwet, Allen P. Kaplan

O-02 C1 Inhibitor function with kinin forming enzymes as target: evaluation of a novel assay
Arije Ghannam, Federica Defendi, Bertrand Favier, Denise Ponard, Christian Drouet

O-03 High Molecular Weight Kininogen, an informative angioedema diagnostic tool
Rémi Baroso, Delphine Charignon, Federica Defendi, Françoise Csopaki, Bertrand Favier and Christian Drouet on behalf of the National Reference Centre for Angioedema CREAK

O-04 Enzymatic assays for the biological diagnosis of BK-dependent angioedema
Federica Defendi, D. Charignon, A. Ghannam, R. Baroso, F. Csopaki, M. Allegret-Cadet, D. Ponard, B. Favier, S. Cichon, B. Nicolie, O. Fain, L. Martin and C. Drouet on behalf of the National Reference Centrefor Angioedema CREAK

O-05 Hormone modulation in women with Hereditary angioedema
Anne Gompel, Isabelle Boccon-Gibod, Laurence Bouillet, David Launay, Ludovic Martin, Gisele Kanny, Olivier Fain

10:25-10:55 **Coffee Break**

10:55-12:55 Scientific lectures *Chairs: Francois Marceau, Andrea Zanichelli*

GENES AND CASCADES

O-06 De novo homozygous mutation of the C1 Inhibitor gene in a patient with hereditary angioedema
V. Bafunno, Chiara Divella, F. Sessa, G.L. Tiscia, G. Castellano, L. Gesualdo, M. Margaglione, V. Montinaro

O-07 Hereditary angioedema due to C1 Inhibitor deficiency among Brazilian patients
Camila Lopes Veronez, Nathália Cagini, Márcia Buzolin, Lício Augusto Velloso, Eli Mansour, João Bosco Pesquero

O-08 Modulation of adrenomedullin and Plaur genes in the pathogenesis of angioedema due to inherited C1 Inhibitor deficiency
Giuseppe Castellano, Chiara Divella, Fabio Sallustio, Vincenzo Montinaro, Andrea Zanichelli, Erika Bonanni, Chiara Suffritti, Loreto Gesualdo, Marco Cicardi

O-09 Gene regulation in the pathogenesis of inherited C1 Inhibitor deficiency (Hereditary Angioedema)
Sonia Caccia, Rosaria Russo, Fleur Bossi, Romina Berardelli, Anna Maria Fra, Valeria Rimoldi, Rosanna Asselta, Stefano Duga, Marco Cicardi

O-10 Immunological method for specific measurement of C1 Inhibitor polymers in plasma and C1 Inhibitor concentrate
Daniel Elenius Madsen, Yaseelan Palarasah, Johannes J. Sidelmann

O-11 Activation of the FXII dependent kallikrein-kinin system by C1-inh polymers
Daniel Elenius Madsen, Johannes J. Sidelmann, Jørgen Gram

O-12 Studies of non-contact activation of Hereditary Angioedema plasma types I, II, and III
Joseph Kusumam, Baby G. Tholanikunnel, Sonia Bains, Konrad Bork, Anette Bygum, Anne Aabom, Claus Koch, Berhane Ghebrehiwet, Allen P. Kaplan

O-13 Inhibition of the plasma kallikrein-kinin system activation by DX-2930, a fully human monoclonal antibody inhibitor of plasma kallikrein
Daniel J. Sexton, Ryan Faucette, Jon Kenniston, Christopher TenHoor, Andrew E. Nixon, Burt A. Adelman

12:55-14:25 **Lunch Break**

14:25-15:55 Poster Session *Chairs: Emel Aygören Pürsün, Timothy Craig*

MIXING IT WITH THE AUTHORS – SCIENCE IN THE SHOP-WINDOW I.

P-01 The Hereditary Angioedema Burden of Illness Study in Europe: A Conceptual Model of Patient Impacts
Anette Bygum

P-02 Hereditary angioedema in Switzerland
Urs Steiner, Werner Pichler, Walter Wuillemin

P-03 Leukocytosis and hemoconcentration yield misinterpretation in abdominal emergencies of Hereditary Angioedema
Daisuke Honda, Isao Ohsawa, Seiji Nagamachi, Hiyori Suzuki, Atsuko Hisada, Satoshi Horikoshi, Yasuhiko Tomino

P-04 Influence of psychological factors on Hereditary Angioedema attacks in children and adolescents: the parents' point of view
Raffaella De Falco, Maria Bova, Maria Francesca Freda, Anna Galante, Angelica Petraroli, Livia Savarese, Gerarda Siani, Massimo Triggiani, Paolo Valerio

P-05 New abdominal crisis trigger on patients with Hereditary Angioedema
José E. Fabiani, Marta Villaverde, Fernando Vazquez Godoy

P-06 No evidence for disease-modifier genes in the PBMC expression profile from HAE patients in basal conditions. An RNA-based microarray screening
Alberto López Lera, Fátima Sánchez Cabo, Sofía Garrido, Ana Dopazo, Margarita López Trascasa

P-07 Gallstones in C1 Inhibitor Deficiency – Are they different?
Jimmy Gooi, Dimitra Athanasiadou, Dimosthenis Sokaras, Udo Becker, Elias Chatzitheodoridis, Athanasios Godelitsas

P-08 The new change in the SERPING1 gene in a Brazilian family can be associated to Hereditary Angioedema (HAE)
Nathália Cagini, Camila Lopes Veronez, Márcia Buzolin, Lício Augusto Velloso, Eli Mansour, João Bosco Pesquero

P-09 Severe laryngeal edema and anemia as first symptoms of IgG kappa multiple myeloma – an acquired angioedema (AAE) case demonstration
Marcin Stobiecki, Krystyna Obtulowicz, Artur Jurczyszyn

P-10 Novel disease severity markers in Hereditary Angioedema due to C1 Inhibitor deficiency
Dorottya Csuka, Lea Munthe-Fog, Mikkel-Ole Skjoedt, Andrea Kocsis, Zsuzsanna Zotter, Péter Gál, Lilian Varga, Henriette Farkas, George Füst†, Peter Garred

P-11 Angioedema - not always an allergic symptom
Kristina Lings, Anette Bygum

P-12 Reduced C1 Inhibitor secretion in fibroblasts derived from patients with Hereditary Angioedema
Iben Rose Christiansen, Claus Koch, Anette Bygum, Jacob Giehm Mikkelsen

P-13 Estrogenic hereditary angioedema
Hakim Rahmoune, Nada Boutrid, Belkacem Bioud

P-14 Possible causes of acquired C1 Inhibitor deficiency, based on our 9 patients
Katalin Molnár, Laura Horváth, Szabolcs Benedek, Lilian Varga, Zsuzsanna Zotter, Henriette Farkas

P-15 Elevated adrenomedullin and endothelin-1 levels during HAE attacks
Erika Kajdácsi, Péter Károly Jani, Dorottya Csuka, Lilian Varga, Zoltán Prohászka, Henriette Farkas, László Cervenak

P-16 Age related changes in the severity of Hereditary Angioedema due to C1 Inhibitor deficiency
Nóra Veszeli, Dorottya Csuka, Lilian Varga, Henriette Farkas

P-17 Adverse effects of danazol prophylaxis in female patients with Hereditary Angioedema due to C1 Inhibitor deficiency (HAE-C1-INH)
Zsuzsanna Zotter, Ibolya Czaller, Dorottya Csuka, Kinga Viktória Kőhalmi, Lilian Varga, Henriette Farkas

P-18 Thyroid function parameters in patients with hereditary angioedema due to C1-inhibitor deficiency
Ibolya Czaller, Dorottya Csuka, Zsuzsanna Zotter, Erika Szabó, Edit Takács, Lilian Varga, Henriette Farkas

P-19 Flow-mediated dilation: assessment of endothelial cell function in patients with hereditary angioedema due to C1-inhibitor deficiency
Zsuzsa Nébenführer, Erika Szabó, Katalin Keltai, András Zsáry, Dorottya Csuka, Zsuzsanna Zotter, Lilian Varga, László Cervenak, Henriette Farkas

15:55-16:25 **Coffee Break**

16:25-17:40 Scientific lectures *Chairs: Coen Maas, Avner Reshef*

THE ATTACK IN THE LIMELIGHT

O-14 Early biomarkers are required for better prediction, evaluation and treatment of HAE attacks
Avner Reshef, Mona Kidon-Iancovici, Boris Gilbord

O-15 Activation of the ficolin-lectin pathway during attacks of Hereditary Angioedema
Dorottya Csuka, Lea-Munthe-Fog, Estrid Hein, Zsuzsanna Zotter, Lilian Varga, Zoltán Prohászka, Peter Garred, Henriette Farkas

O-16 A preliminary study into the activation of plasma enzyme systems during attacks of Hereditary Angioedema due to C1 Inhibitor deficiency (HAE-C1-INH)
Dorottya Csuka, Éva Imreh, Zsuzsanna Zotter, Szilvia Walentin, Mónika Kleiber, Lilian Varga, Henriette Farkas

O-17 Endothelial cell activation during edematous attacks of Hereditary Angioedema type I and II
Erika Kajdácsi, Péter Károly Jani, Dorottya Csuka, Lilian Ágnes Varga, Zoltán Prohászka, Henriette Farkas, László Cervenak

O-18 Activation of the Plasma Contact-system in patients with Anaphylaxis
A. Sala Cunill, M. Guilarte, V. Cardona, J. Björkqvist, M. Labrador, K. Nickel, T. Renné

19:15- **Social dinner**

Saturday, 25 May

08:30-09:00 **Prize-winner lecture and Distribution of prize „for HAE Research Award"**
Chair: Bruce Zuraw
Awarded: *Allen Kaplan*

09:00-10:45 Scientific lectures *Chairs: Janne Björkander, Konrad Bork*

SORTING THINGS OUT: HAE-FXII

O-19 Hereditary angioedema with normal C1 Inhibitor: Clinical characteristics of seven families in Catalonia
Maria del mar Guilarte, A. Sala-Cunill, M. Labrador-Horrillo, O. Luengo, V. Cardona

O-20 A deletion in the factor 12 gene analysed in two Turkish families with Hereditary Angioedema with normal C1 Inhibitor (HAE type III): a Turkish *F12* mutant?
Konrad Bork, Karin Wulff, Jochen Hardt, Guenther Witzke

O-21 Mutation in Coagulation Factor XII gene associated with Hereditary Angioedema with normal C1 Inhibitor in Brazilian families
Adriana S. Moreno, S.O.R. Valle, A.T. França, S.A. Levy, F.S. Serpa, N. Monnier, D. Ponard, J. Lunardi, M.D. Mendonça, W.N. Campos, H. Arcuri, M.S. Palma, W.A. Silva Junior, L.K. Arruda

O-22 Hereditary Angioedema-factor XII associated. Clinical and genetic characteristics in an Andalusian cohort (South of Spain)
Macarena Piñero-Saavedra, T. González-Quevedo, R. García-Lozano, B. Saenz de San Pedro

O-23 Hereditary Angioedema patients with normal C1 Inhibitor and factor XII mutation: a French cohort
Isabelle Boccon-Gibod, Anne Gompel, Olivier Fain, Yann Ollivier, Nadia Raison-Peyron, Aurélie Du-Than, Stéphane Gayet, Laurence Bouillet

O-24 Hereditary Angioedema with factor XII mutation: no evidence for contact activation during attack (case report)
Chiara Suffritti, Andrea Zanichelli, Marta Mansi, Christiane Stieber, Giulia Periti, Erika Bonanni, Lorena Maggioni, Romualdo Vacchini, Marco Cicardi

O-25 Kinin catabolism and disease severity in hereditary angioedema with *F12* mutation
Delphine Charignon, Federica Defendi, Arije Ghannam, Denise Ponard, Sven Cichon, Olivier Fain, Ludovic Martin, Christian Drouet on behalf of the CREAK, the National Reference Centre for Angioedema

10:45-11:15 **Coffee Break**

11:15-13:00 Scientific lectures *Chairs: Jonathan Bernstein, Grzegorz Porebski*

MANAGEMENT FINE-TUNED

O-26 Long term prophylactic treatment in Spanish patients with Hereditary Angioedema with C1 Inhibitor Deficiency (HAE-C1INH)
Maria Pedrosa, M. Guilarte, T. González Quevedo, M.L. Baeza, T. Lobera, C. Marcos, B. Sáenz de San Pedro, J. Jurado, T. Caballero

O-27 Anabolic androgen experience and response to nanofiltered C1 inhibitor from the CINRYZE prevention trials in patients with HAE
B.L. Zuraw, D. Mariano, J. Dayno

O-28 HAE Patient experience with short term prophylaxis: Responses to an on-line questionnaire survey
Jonathan A. Bernstein, Umesh Singh, Joyce Wilmot

O-29 Management of Hereditary Angioedema. Real-world experiences from a Danish Specialist Centre
Anette Bygum

O-30 Prospective evaluation of the efficacy of on demand treatments in reducing duration of angioedema in patients with Hereditary Angioedema due to C1 inhibitor deficiency
Giulia Periti, Andrea Zanichelli, Marta Mansi, Romualdo Vacchini, Chiara Suffritti, Lorena Maggioni, Erika Bonanni, Marco Cicardi

O-31 Ruconest in routine clinical practice: UK experience
A.L. Manson, J. Dempster, S. Grigoriadou, M.S. Buckland, Hilary J. Longhurst

O-32 Clinical trial experience of pediatric patients treated with ecallantide for acute attacks of Hereditary Angioedema
Andrew J. MacGinnitie, Mark Davis-Lorton, Leslie E. Stolz, Raffi Tachdjian, Ibrahim Dagher

13:00-14:30 **Lunch Break**

14:30-16:00 Poster Session *Chairs: Anette Bygum, Tom Bowen*

MIXING IT WITH THE AUTHORS – SCIENCE IN THE SHOP-WINDOW II.

P-20 A cross-sectional questionnaire survey to assess physician's approach to short-term prophylaxis in HAE patients
Jonathan A. Bernstein, Umesh Singh, Joyce Wilmot

P-21 Case report of use of Icatibant during pregnancy
Andrea Zanichelli, Marta Mansi, Giulia Periti, Chiara Suffritti, Lorena Maggioni, Erika Bonanni, Romualdo Vacchini, Marco Cicardi

P-22 Experience and current status of the translation and cross-cultural adaptation of the angioedema quality of life questionnaire (AE-QoL)
Karstem Weller, Marcus Maurer, Markus Magerl

P-23 The Romanian Hereditary Angioedema Registry
Dumitru Moldovan, Eniko Mihaly, Noemi Anna Bara, Valentin Nădăşan

P-24 Pilot study and validation of the IHAE-QoL questionnaire
Nieves Prior, E. Remor, E. Pérez-Fernández, C. Gómez-Traseira, M. Caminoa, F. Gayá, A. Aabom, W. Aberer, S. Betschel, A. Bygum, D. Csuka, H. Farkas, A. Groffik, M. Gomide, A. Grumach, I. Leivobich, A. Malbran, E. Mihaly, D. Moldovan, K. Obtulowicz, G. Porebski, C. Rayonne, A. Reshef, P. Staubach, M. Wiednig, T. Caballero

P-25 The US HAE Association: an important partner in a longstanding effort to improve patients' quality of life
Anthony Castaldo, J.F. Long, D.K. Davis, L.I. Perry, D.L. Williamson, P.L. King

P-26 Hereditary Angioedema - how can medicines reach the patient?
Maja Jošt, Mihaela Zidarn, Mitja Košnik

P-27 Comparison of acute angioedema attacks versus breakthrough attacks during a placebo-controlled, crossover study of CINRYZE® (C1 Esterase Inhibitor [Human]) for prophylaxis in patients with Hereditary Angioedema
Jennifer Schranz, D. Fitts, C. Broom

P-28 Feasibility of home infusion and self-administration of CINRYZE® (C1 Esterase Inhibitor [Human]) for routine prophylaxis in patients with Hereditary Angioedema and characterization of a training and support program
David Mariano, L.M. Landmesser, C. Gregory

P-29 Clinical descriptive study and Health Related Quality of Life (HRQoL) as measured by SF-36v2 in adults with Hereditary Angioedema due to C1-inhibitor deficiency in Spain
Teresa Caballero, C. Gómez-Traseira, M. Caminoa, E. Pérez-Fernández, C. Andreu, A. Campos, P. Carretero, L. Fernández-Vieira, A. Ferrer, F. García-González, T. González-Quevedo, M. Guilarte, M.A. Gonzalo-Garijo, C.H. Larramendi, T. Lobera, C. Marcos, A. Salas, P. Sánchez-Payá, M.E. Sanchís, M.T. Soto-Mera, N. Prior

P-30 Benefits of early administration of Icatibant for the treatment of Hereditary Angioedema attacks
Hilary Longhurst, Werner Aberer, Laurence Bouillet, Teresa Caballero, Vincent Fabien, Priscila Valente de Freitas, Andrea Zanichelli, Marcus Maurer

P-31 Treatment of Hereditary Angioedema attacks with Icatibant: A comparison of observational data with clinical trial data
Marcus Maurer, Hilary Longhurst, Michaela Wiednig, Vincent Fabien, William Lumry

P-32 Teaching intravenous self-application in patients with HAE: Experiences by a specialist nurse
H. Mühlberg, Nicole Ettl, M. Magerl

P-33 Different forms of HAE prophylaxis
Murat Bas, Ulrich Straßen

P-34 An indirect comparison of icatibant and four other therapies for the symptomatic treatment of acute attacks of Hereditary Angioedema types I and II
Matthew Helbert, M. Alvarez-Reyes, I. Pearson, L. Diwakar

P-35 The efficacy and safety of self-injected icatibant administered as an acute treatment for Hereditary Angioedema due to C1 Inhibitor deficiency (HAE-C1-INH) in clinical practice
Zsuzsanna Zotter, Dorottya Csuka, Lilian Varga, György Temesszentandrási, Henriette Farkas

P-36 Short-term prophylaxis in a patient with acquired C1-INH deficiency
Kinga V. Kőhalmi, Zsuzsanna Zotter, Dorottya Csuka, Katalin Molnár, Lilian Varga, Henriette Farkas

P-37 Home treatment of attacks with conestat alfa in Hereditary Angioedema due to C1 Inhibitor deficiency (HAE-C1-INH)
Erika Szabó, Dorottya Csuka, Zsuzsanna Zotter, Lilian Varga, Henriette Farkas

16:00-16:30 **Coffee Break**

16:30-17:45 Scientific lectures *Chairs: Laurence Bouillet, Hilary Longhurst*

ANGIOEDEMA AVALANCHE

O-33 Follow-up of patients with drugs targeting the renin-angiotensin-aldosterone system-induced angioedema
Macarena Pinero-Saavedra, Isabelle Boccon-Gibod, Teresa Gonzalez-Quevedo, Laurence Bouillet

O-34 Asphyxiation in HAE due to C1-INH deficiency and HAE with normal C1-INH
Konrad Bork, Jochen Hardt, Günther Witzke

O-35 Clinical survey of different forms of angioedema without wheals
Marta Mansi, Andrea Zanichelli, Giulia Periti, Chiara Suffritti, Erika Bonanni, Lorena Maggioni, Romualdo Vacchini, Marco Cicardi

O-36 Surveys of prodromes preceding acute attacks of Hereditary Angioedema
Avner Reshef, Michael J Prematta, Timothy J. Craig

O-37 The influence of age at first clinical manifestation of Hereditary Angioedema (HAE) on the clinical course of the disease
Inmaculada Martinez-Saguer, Eva Rusicke, Carmen Escuriola-Ettingshausen, Emel Aygören-Pürsün, Karin Andritschke, Adrianna Piotrowski, Wolfhart Kreuz

18:30- **Social dinner**

Sunday, 26 May

08:30-09:45 Scientific lectures *Chairs: Iris Leibovich, Alejandra Menendez, Dumitru Moldovan*
THE SHADES OF FREEDOM – Nursing round table discussion

O-38 Living with Hereditary Angioedema – Nursing aspects
Iris Leibovich

O-39 Practical Approach to self-administration of C1-Inhibitor in HAE-Patients
Karin Andritschke

O-40 The Hungarian Patient – Hospitalisation of HAE-patients with diseases different from HAE
Arianna Kitzinger

O-41 Practicalities and barriers to HAE self-administration therapy: discussions from an international HAE expert meeting
Christine Symons, H.B. Boysen, L. Bouillet, S. Neri, J. Hébert, E. Aygören-Pürsün, I. Martinez-Saguer, C. Bethune, A. Sala-Cunill, M. Cancian, O. Rossi, M. Magerl, K. Andritschke, T. Craig

O-42 The building blocks for an effective patient group advocacy program
Anthony Castaldo, H.B. Boysen, M. Rutkowski, S.F. Smith, A. Menendez, H. Mykal, J. Schultz-Boysen, V. Ledbez, P. Hermeling

09:45-10:15 **Coffee Break**

10:15-12:30 Scientific lectures *Chairs: Teresa Caballero, Bernd Rosenkranz*
LIVING WITH HAE AROUND THE WORLD

O-43 International clinical descriptive study of adults with Hereditary Angioedema due to C1 Inhibitor deficiency
Teresa Caballero, M. Caminoa, E. Pérez-Fernández, C. Gómez-Traseira, F. Gayá, A. Aabom, W. Aberer, S. Betschel, A. Bygum, D. Csuka, H. Farkas, A. Groffik, M. Gomide, A. Grumach, I. Leivobich, A. Malbran, E. Mihaly, D. Moldovan, K. Obtulowicz, G. Porebski, C. Rayonne, A. Reshef, P. Staubach, M. Wiednig, N. Prior

O-44 Clinical differences among countries in Hereditary Angioedema due to C1 Inhibitor deficiency
Nieves Prior, Magdalena Caminoa, Elia Pérez-Fernández, Carmen Gómez-Traseira, F. Gayá, Anne Aabom, Werner Aberer, Stephan Betschel, Anette Bygum, Dorottya Csuka, Henriette Farkas, Adriane Groffik, M. Gomide, Anete Grumach, Iris Leivobich, A. Malbran, Eniko Mihaly, Dumitru Moldovan, K. Obtulowicz, Grzegorz Porebski, C. Rayonne, Avner Reshef, P. Staubach, Michaela Wiednig, Teresa Caballero

O-45 Health status utility weights for Hereditary Angioedema attacks and in between attacks
Emel Aygören-Pürsün, Teresa Caballero, Anette Bygum, Kathleen Beusterien, Emily Hautamaki, Zlatko Sisic, Suzanne Wait, Henrik B. Boysen

O-46 HAE – The Situation in South Africa
Bernd Rosenkranz

O-47 Hereditary Angioedema in Latin America: 1st report
J. Fabiani, S.O.R. Valle, M. Olivares, S. Nieto, E.H. Landeros, A. Ginaca, L. Bezrodnik, E. Nievas, M. Oleastro, O.M. Barrera, A.M. Gallardo, A. King, J.R. Galindo, M.J.O. Carabantes, M.M.Alfonso, R. Vilarim, Anete S. Grumach

O-48 Quality of life and productivity loss in patients with Hereditary Angioedema (HAE) in Sweden; results from a retrospective patient registry survey implemented by Sweha-eg (A population based census of HAE in Sweden)
Patrik Nordenfelt, J. Björkander, L. Mallbris, A. Lindfors, S. Friberg, K. Löfdal, L. Nordvall, S. Werner, C.F. Wahlgren

O-49 Hereditary Angioedema in Greece
F.E. Psarros, M. Speletas, N. Koutsostathis, Anastasios E. Germenis

O-50 HAE in Macedonia: Current status
Vesna Grivcheva-Panovska

O-51 Hereditary Angioedema nationwide study in Slovenia
Matija Rijavec, Peter Korošec, Mira Šilar, Mihaela Zidarn, Jovan Miljković, Mitja Košnik

12:30-13:00 **Closing Ceremony** *Chairs: Henriette Farkas, Peter Späth*
Distribution of prizes for „Grant for Young Investigators"
Closing remarks by Peter Späth

13:00-14:30 **Lunch**

14:30- **Farewell & Departure**